Limberg Trial

No registrations found.

Ethical review	Not applicable
Status	Pending
Health condition type	-
Study type	Interventional

Summary

ID

NL-OMON22628

Source
Nationaal Trial Register

Brief title
Limberg Trial

Intervention

Outcome measures

Primary outcome

Tijd tot volledige wondgenezing

Secondary outcome

Het aantal wondinfecties, het recidiefpercentage binnen 1 jaar, en een aantal factoren die de morbiditeit na de operatie bepalen zullen worden onderzocht.

Study description

Study design

n.v.t.

Intervention

Limbergplastiek vs. sluiten in midline na excisie sinus pilonidalis

Contacts

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Eligibility criteria

Inclusion criteria

primary or recurrent pilonidal sinus
width of fistulacomplex between 1 and 3 centimeter
signed informed consent
age: >/= 18 years
mentally competent

Exclusion criteria

patients with an actual abscess
not able to follow-up
patients in which woundhealing problems are to be expected (Diabetes type II, immunesupression, chemotherapy, HIV/AIDS)
not mentally competent
dementia

Study design

Design

Study type:	Interventional
Intervention model:	Parallel
Allocation:	Randomized controlled trial
Masking:	Single blinded (masking used)
Control:	Active

Recruitment

NL

Recruitment status:	Pending
Start date (anticipated):	01-01-2015
Enrollment:	74
Type:	Anticipated

Ethics review

Not applicable

Application type:	Not applicable

Study registrations

Followed up by the following (possibly more current) registration

ID: 44556
Bron: ToetsingOnline
Titel:

Other (possibly less up-to-date) registrations in this register

No registrations found.

In other registers

Register	ID
NTR-new	NL4792
NTR-old	NTR4932
CCMO	NL49989.044.14
OMON	NL-OMON44556

Study results